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(54) Intumescent material

(57) An intumescent material comprises: one or more organic polyhydroxy compounds as carbonific; one or more organic polyamido compounds in free and/or combined form as spumific; an ammonium phosphate as activator for the carbonific; and an amino resin as binder, the material further comprising sodium silicate in an amount insufficient to prevent the material intumescing on being subjected to elevated temperature.

A method of manufacturing the intumescent material, comprises: preparing an aqueous paste or slurry comprising the ingredients of the material and allowing or causing the paste or slurry to harden.

SPECIFICATION

Intumesc nt mat rial

	and the second man sign	5
5	The pr sent invention relates to intumescent mat rial. An intumescent material is a material which when heated or subjected to elevated tempera-	
	to the conditions expands to form 8 SOUG TOSMI.	
	a i i i i i i i i i i i i i i i i i i i	
		10
10		10
	A second known type of intumescent material comprises components acting as carbonius,	
	10	
	The carbonific is a source of carbon char when the intumescent material is subjected to heat. It is the principal constituent of the foam formed on intumescence. Mono-, di- and tri-	15
15		
	The annual decomposes under tire conditions and releases usses which decode the decode.	
	char to expand. Polyamido compounds such as dicyandiamide and guanidine, in free or	
	4: 1 Company and averaging of OFGRADIC SPIRMITICS	20
20	The activator promotes the decomposition of the carbonitic. Antificituding prospilates dusing a	20
20	nhochate are examples of activators.	
	The binder may be an amino resin such as urea-tormaidenyde resin or melanime-tormain	
	The ansign also gots to some extent as a carponitic and a spulling.	
	Thus the second known type of intumescent material may comprise an amino resin as binder	25
25	and an ammonium phosphate such as monoammonium phosphate as activator, together with carbonific and spumific. Such a material needs to be formulated so as to exhibit an acidic	
	ammonium phosphate has to yield phosphoric acid in order to promote the decomposition of the	
	, ,,,,	
30	taken and motorials of the second type and comprising amino resin as billider are discussed	30
50		
	debute resignators linked by a polyamido compound, the cross-linked resin acting as billed and	
	the polyamide compound acting as spumitic. The material also comprises all altitionism	
	think are an entire to a property polyhydroxy compound as carponiic.	35
35	Intumescent materials, as described above, made from cross-linked amino resins, such as	•
	melamine-formaldehyde resin, may be regarded as heavily filled amino resins and hence are stronger and less friable than sodium-silicate based intumescent material.	
	Francisco entirestant interpretations interpretations materials based on amino resins, especially inclaiming	
	formaldehyde resin may be considered superior to intumescent materials comprising sociality	
40	alliants in view of the shove-mentioned differences in their properties.	40
40	the second second control of the second control of the second sec	
	formaldehyde resin and hence intumescent materials based on sodium silicate can be manufac-	
	are detailed cost than intumescent materials based on amino resins.	
	The above described intumescent materials comprising amino resins are generally manufac-	45
45	tured by making a thick aqueous slurry or paste of the required ingredients and allowing or	
	causing the slurry or paste to harden. In the case of intumescent materials as described in our aforementioned British patent, the	
	Abiels agreed a clurpy or paste is prepared from inter alla Water-dispersible melamille-lumina-	
	dehyde resin, a polyamido compound such as dicyandiamide as cross-linking agent for the resin	
50	and an assumition and managementum phosphate.	50
•	It would not be thought that a satisfactory incumescent material comprising components	
	period as earbonific, solumific and binder, ammonium phosphate acting as activator, and also	
	andium cilicate, could be manufactured at reduced cost (as compared with such a material	
	without sodium silicate) because the alkalinity of the sodium silicate would be expected to	55
55	prevent or hinder the yielding of phosphoric acid by the ammonium phosphate. Also in manufacturing the intumescent material, if sodium silicate were incorporated into the aqueous	
	slurry or paste, it would be expected to react with the resin, which gives an acidic reaction in	
	weeter, and provent the resin cross-linking properly.	
	We have now surprisingly discovered that a satisfactory and, in some respects superior,	_
60	intumescent material may comprise a carbonitic, an organic spumitic, an amino resit, such as	60
55	areas linked melamine-formaldehyde resin, as binder, and sodium silicate.	
	Accordingly the present invention provides an intumescent material comprising; one or more	
	accomic polybydroxy compounds as carbonitic; one or more organic polyamido compounds in	
	to and/or combined form as snumific an ammonium phosphate as activator for the carbonnic,	65
65	and an amino resin as binder, the material further comprising sodium silicate in an amount	٠.

insufficient to pr vent the material intumescing on being subjected to I vated t mp rature. It will be understood that the sodium silicat in a sinse constitutes a further spumific. The proportions of the organic polyhydroxy compound, the polyamido compound, the amm nium phosphate, the amino r sin, and any other ingredi nts (except sodium silicate) may 5 5 be varied as describ d in our British patent no. 1601131. The amino resin is preferably melamine-formaldehyde resin cross-linked by a polyamido compound, such as dicyandiamide or guanidine, which acts as spumific. The carbonific may conveniently be pentaerythritol. The activator is preferably monoammonium phosphate or may be ammonium polyphosphate. In order to provide satisfactory intumescence, the amount of sodium silicate preferably does 10 not exceed 12.7%, more preferably 11.5% by weight based on the total weight of the aforementioned carbonific, organic spumific (polyamido compound), activator and binder. Preferably the amount of sodium silicate does not exceed 32.6%, more preferably 29.6%, by weight based on the ammonium phosphate. 15 Preferably the amount of sodium silicate does not exceed 36%, more preferably 32.8%, by weight based on the amino resin excluding any cross-linkages thereof (e.g. cross-linking polyamido compound where the resin is melamine-formaldehye resin cross-linked by such polyamido compound). Preferably the amount of sodium silicate does not exceed 63.5%, more preferably 62.3%, by 20 20 weight based on the polyamido compound. Preferably the amount of sodium silicate does not exceed 171%, more preferably 155%, by weight based on the polyhydroxy compound. Preferably the amount of sodium silicate does not exceed 11%, more preferably 10%, by weight based on the ingredients of the material other than sodium silicate and any water 25 25 present. In order to provide a stiff "puff" as described below, the amount of sodium silicate preferably is at least 3.46%, more preferably 4.6%, by weight based on the total weight of the aforementioned carbonific, spumific (polyamido compound), activator and binder. Preferably the amount of sodium silicate is at least 8.9%, more preferably 11.9%, by weight 30 30 based on the ammonium phosphate. Preferably the amount of sodium silicate is at least 9.8%, more preferably 13.1% by weight based on on the amino resin excluding any cross-linkages thereof (e.g. cross-linking polyamido compound where the resin is melamine-formaldehyde resin cross-linked by such polyamido compound). Preferably the amount of sodium silicate is at least 18.7%, more preferably 24.9%, by weight based on the polyamido compound. Preferably the amount of sodium silicate is at least 46.7%, more preferably 62.3%, by weight based on the polyhydroxy compound. Preferably the amount of sodium silicate is at least 3%, more preferably 4%, by weight based 40 40 on the ingredients of the material other than sodium silicate and any water present. The intumescent material of the invention may further comprise polyvinyl acetate as additional binder to provide with the amino resin an interpenetrating polymer network system, thereby making the intumescent material less brittle. The intumescent material of the invention may further comprise a hardened setting agent such as hardened plaster of Paris. The intumescent material according to the invention has a number of advantages, apart from 45 its reduced cost, as compared with the second known type of intumescent material. On heating intumescent material according to the invention, intumescence occurs in two stages. A first stage of intumescence occurs at a relatively low temperature due to the presence of the sodium silicate and the second stage of intumescence occurs at a somewhat higher 50 50 temperature due to the ingredients of the second type of intumescent material. Moreover, the second stage intumescence tends to occur more slowly than if the sodium silicate were absent. Thus although the first stage intumescence may occur quickly, the second stage intumescence may be advantageously prolonged. Moreover the intumesced material or "puff" resulting from heating of intumescent material 55 according to the invention is stiff. In contrast the "puff" obtained from the conventional sodium 55 silicate-based intumescent material is hard and the "puff" obtained from intumescent material of the second type is soft and elastic. A stiff "puff" may be advantageous in certain situations for example in retarding warping of doors under fire conditions. Under such conditions a soft and elastic "puff" would deform easily 60 60 to accommodate the warping of the door and may slump to leave a gap between the intumesced material and the door whereas a hard "puff" would crack and crumble and may tend to force th door op n. However, a stiff "puff" has sufficient rigidity not to slump but do s not t nd to crack and crumble r to force the door open. Mor over, when the intumescent material according to the invention intumesces through an 65 65 op ning in a hold r it tends to expand in a direction normal to the opening rather than

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spreading laterally from the opening and can thus seal larger gaps than the known intumescent materials. The degree of "puff" or intumescence of the intumescent material may be up to 1/3 as large again as that of known intumescent materials of the second type and as described in our 5 aforementioned British patent. We have also found that the intumescent material according to the invention is not adversely affected by prolonged exposure to high concentrations of carbon dioxide or atmospheric moisture. The material can withstand 100% relative humidity without its intumescent properties being adversely affected. We have furthermore found that the intumescent material according to the invention retains 10 its intumescent potential well under accelerated ageing even for ageing equated with as much as ten years. The intumescent material of the invention may be manufactured by preparing an aqueous paste or slurry comprising: one or more organic polyhydroxy compounds as carbonific, one or 15 more polyamido compounds as spumific, an ammonium phosphate as activator and an amino 15 resin as binder, and sodium silicate, and allowing or causing the paste or slurry to harden, the amount of sodium silicate being insufficient to prevent the material manufactured intumescing on being subjected to elevated temperature. Preferably the amino resin is water-dispersible melamine-formaldehyde resin, which becomes 20 cross-linked by the polyamido compound. The polyamido compound may be dicyandiamide or 20 guanidine, which can effect cross-linking of melamine-formaldehyde resin at normal ambient temperature (15 to 25°C). The amount of sodium silicate in relation to the other ingredients of the material is preferably as specified above in relation to the intumescent material of the invention. The aqueous paste or slurry may further comprise water-dispersible polyvinyl acetate as binder 25 (in addition to the amino resin) for the purpose referred to above and a setting agent such as plaster of Paris. Preferably the pH of the paste or slurry from which the intumescent material is prepared is not above 6.2, more preferably 6.1, otherwise the degree of intumescence of the material on 30 heating may be inadequate. 30 We have found that the paste or slurry runs very smoothly and is thus advantageous for loading into holders in which it subsequently sets to form intumescent material. Also the paste or slurry tends to be less adherent to the equipment used in its preparation than the aqueous mixture used in preparing the known intumescent materials and thus leaves the equipment 35 cleaner. Preferably the pH of the paste or slurry is about 6.15 to provide an optimum 35 combination of smoothness of the paste or slurry and adequate intumescence of the intumescent material, although a lower pH would normally be chosen in order to provide greater intumescence. The invention is illustrated by the following Examples: The sodium silicate used in the Examples was commercially available sodium silicate having 40 an SiO: Na2O molar ratio of 1:2. The plaster of Paris used in the Examples was of a kind which exhibits high expansion during setting. Example 1 45 A powder mix of the following ingredients in the stated amounts was prepared:-45 water-dispersible melamine formaldehyde 1900g resin powder monoammonium phosphate 2100g 50 dicyandiamide 1000g 50 400g pentaerythritol plaster of Paris 550g 150g water-dispersible polyvinyl acetate powder wood flour (passing through 90 mesh) 125g

100 parts by weight of this powder mix were mixed with various amounts of sodium silicate powder, the resulting mixture worked up with water and the pH of the worked-up mixture 60 measured. The worked-up mixture was then allowed to set and dry. The set and dried mixture was then subject to elevated temperature to determine its degree of intumescence.

Total 6225g

The results were as follows:

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5 WE	.licate (parts b eight)	v PH	Intumescence	Remarks		5		
.	. _ 3 ,	-						
	4	< 6	good	stiff "puff" o				
0	6	< 6	very good	stiffer "	••	10		
	8	~ 6	11 11	stiffest "	11			
	10	6.1	good					
5	12½	6.5	reduced			15		
	15	6.8	poor					
20	xample 2 The following comp	position was made	as a dry powder			20		
m di 25 pi pi w	ater-dispersible mela conoammonium pho icyandiamide entaerythritol laster of Peris rater-dispersible poly	sphate	2100g 1000g 400g 550g			25		
	sodium silicate powder 498g The amount of sodium silicate was thus 8% by weight based on the other ingredients.							
S1	Water was mixed with the above dry composition to form a thick aqueous slurry. The thick aqueous slurry was quickly poured into a tubular holder strip. The initial setting of the material started within 1-2 hours and chemical setting was well advanced after 2 days. The holder strip then slit along one face by a milling cutter and the holder with the hardened mixture was then kept at 40°C for 24 hours to mature the hardened mixture (i.e. to allow free water to evaporate and to complete the chemical setting). The holder containing the intumescent material is							
k a	ept at 40°C for 24 I nd to complete the	hours to mature th chemical setting).	tter and the holder wi ne hardened mixture (i The holder containing	th the hardened mi .e. to allow free wa	xture was then ter to evaporate	3		
k a rı	ept at 40°C for 24 lend to complete the eferred to herein as A second sealing decision silicate in presenting the present the	hours to mature the chemical setting). a first sealing devi levice was made it poaring the aqueou	tter and the holder wi te hardened mixture (i The holder containing ice. n like manner to that (th the hardened mi .e. to allow free wa the intumescent m described above bu	xture was then ter to evaporate naterial is t using no	3		
k a r 10 s s s \$5 p	ept at 40°C for 24 lend to complete the offerred to herein as A second sealing dodium silicate in presented that the stantially identicate on heating the two odium silicate intum silicat	hours to mature the chemical setting). a first sealing device was made in a paring the aqueous the crack resistance in sealing devices, in a good to be sealing devices, in a device (and colors of the colors of th	tter and the holder with hardened mixture (in the holder containing ice. In like manner to that our slurry. I of the intumescent rether material therein in the more than the other momerising sodium silice.	th the hardened mi .e. to allow free way the intumescent m described above bu material in the two atumesced, the material. The intume	xture was then ter to evaporate naterial is t using no holders was erial made using	4		
40 s 40 s 45 p tl	ept at 40°C for 24 lend to complete the eferred to herein as A second sealing dodium silicate in present the second sealing dodium silicate intum rovided by first sealing doctates. An intumescer arbonific; one or monumitic: an ammonity and to complete the second sealing doctates.	hours to mature the chemical setting). a first sealing device was made in the crack resistance it. The control of the crack resistance it. The crack re	tter and the holder with the hardened mixture (in the holder containing tice. In like manner to that the survey. In the material therein in the material therein in the other material	th the hardened mile. to allow free way the intumescent medescribed above but material in the two fatumesced, the material. The intume ate) was stiff and the material of the	erial made using esced mass nat provided by			
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8.	A material	according	to any	preceding	claim,	wher is	n the	amount of	sodium	silicate o	does
not ex	ceed 11%	by weight	based	n the tota	al weigt	nt of the	e othe	r ingredie	nts of th	materia	ıl.
٥	A material	according	to any	prec ding	claim.	wh rein	n the :	amount of	s dium	silicate i	s at

least 3.46% by weight based on the total weight of the organic polyhydroxy compound, the polyamido compound, the ammonium phosphate and the amino resin.

10. A material according to any preceding claim, wherein the amount of sodium silicate is at least 8.9% by weight based on the ammonium phosphate.

11. A material according to any preceding claim, wherein the amount of sodium silicate is at least 9.8% by weight based on the amino resin excluding any cross-linkages thereof.

12. A material according to any preceding claim, wherein the amount of sodium silicate is at least 18.7% by weight based on the polyamino compound.

13. A material according to any preceding claim, wherein the amount of sodium silicate is at least 46.7% by weight based on the polyhydroxy compound.

14. A material according to any preceding claim, wherein the amount of sodium silicate is at 15 least 3% by weight based on the total weight of the other ingredients of the material.

15. A method of manufacturing an intumescent material, comprising: preparing an aqueous paste or slurry comprising one or more organic polyhydroxy compounds as carbonific, one or more polyamido compounds as spumific, an ammonium phosphate as activator and an amino resin as binder, and sodium silicate, and allowing or causing the paste or slurry to harden, the 20 amount of sodium silicate being insufficient to prevent the material manufactured intumescing

amount of sodium silicate being insufficient to prevent the material manufactured intumescing on being subjected to elevated temperature.

16. A method according to claim 15, wherein the amino resin is water-dispersible melamine-formaldehyde resin and the aqueous paste or slurry comprises a polyamido compound as spumific and cross-linking agent for the resin.

25 17. A method according to claims 15 or 16, wherein the pH of the paste or slurry is not above 6.2.

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